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NEWS 20 JAN 28 USGENE now provides USPTO sequence data within 3 days
of publication
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NEWS 22 JAN 28 MEDLINE and LMEDLINE reloaded with enhancements
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L1 6 (ALPHA 1-ANTITRYPSIN) (6A) POWDER

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=> d l2 1-6 bib ab

L2 ANSWER 1 OF 6 CAPLUS COPYRIGHT 2008 ACS on STN
AN 2007:672983 CAPLUS
DN 147:102152
TI pharmaceutical powder compositions for inhalation
IN Mueller-Walz, Rudi
PA Jagotec A.-G., Switz.
SO PCT Int. Appl., 30pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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PI	WO 2007068443	A1	20070621	WO 2006-EPI1941	20061212
	W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, VZ, VC, VN, ZA, ZM, ZW			
	RW:	AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM			

PRAI GB 2005-25254 A 20051212

AB A pharmacol. powder for inhalation comprising fine particles of a drug and particles of a force-controlling agent, wherein the particles of the force-controlling agent are disposed on the surface of the active particles as either a particulate coating, or as a continuous or discontinuous film. The powder may further comprise particles of a carrier material for supporting the drug particles. The force-controlling agent may be selected from: amino acids, peptides and polypeptides having a mol. weight of 0.25 to 1000 KDa; phospholipids; titanium dioxide; aluminum dioxide; silicon dioxide; starch; and salts of fatty acids. Also disclosed is a method of making such a powder for inhalation comprising mixing a force-controlling agent with particles of one or more pharmacol. active materials to obtain a mixture in which the particles of the force-controlling agent are disposed on the surface of the active particles as either a particulate coating, or as a continuous or discontinuous film. The mixing step may be achieved by sieving, mixing or blending, micronizing and/or co-micronizing the particles of one or more pharmacol. active materials and particles of force-controlling agents. A powder formulation consisting of glycopyrrolate, magnesium stearate and lactose monohydrate was obtained. The dry powder blend achieved is homogeneous and the blend has satisfying blend homogeneity.

RE.CNT 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L2 ANSWER 2 OF 6 CAPLUS COPYRIGHT 2008 ACS on STN
AN 2005:451414 CAPLUS
DN 142:487376
TI Dry recombinant human alpha 1-antitrypsin formulation
IN Nayar, Rajiv; Manning, Mark G.; Barr, Philip J.; Pemberton, Philip A.; Bathurst, Ian C.; Gibson, Helen
PA Arriva-Prometic Inc., Can.
SO PCT Int. Appl., 21 pp.
CODEN: P1XXD2
DT Patent
LA English
FAN.CNT 2

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2005047323	A1	20050526	WO 2004-GB4740	20041110
	W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, VZ, VC, VN, YU, ZA, ZM, ZW			
	RW:	BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK,			

EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LU, MC, NL, PL, PT, RO,
SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR,
NE, SN, TD, TG

AU 2004288854 A1 20050526 AU 2004-288854 20041110
CA 2545458 A1 20050526 CA 2004-2545458 20041110
EP 1685160 A1 20060802 EP 2004-798463 20041110

R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
IE, SI, FI, RO, CY, TR, BG, CZ, EE, HU, PL, SK, IS

JP 2007534633 T 20071129 JP 2006-538939 20041110
US 20070105768 A1 20070510 US 2006-578692 20060826

PRAI US 2003-518803P P 20031110
US 2003-519946P P 20031114
WO 2004-GB4740 W 20041110

AB A dry powder composition comprises recombinant human alpha
1-antitrypsin (rAAAT).

RE.CNT 5 THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L2 ANSWER 3 OF 6 CAPLUS COPYRIGHT 2008 ACS on STN
AN 2001:338322 CAPLUS
DN 134:357557
TI Dry powder compositions having improved dispersivity
IN Lechuga-Ballesteros, David; Kuo, Mei-Chang
PA Inhale Therapeutic Systems, Inc., USA
SO PCT Int. Appl., 56 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2001032144	A1	20010510	WO 2000-US9785	20000412
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW				
	RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
	CA 2389219 A1 20010510		CA 2000-2389219		20000412
	EP 1223915 A1 20020724		EP 2000-922117		20000412
	EP 1223915 B1 20051221				
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL				
	JP 2003513031 T 20030408		JP 2001-534349		20000412
	HU 2003001851 A2 20030929		HU 2003-1851		20000412
	HU 2003001851 A3 20060728				
	NZ 518401 A 20040130		NZ 2000-518401		20000412
	AU 775565 B2 20040805		AU 2000-42353		20000412
	AT 313318 T 20060115		AT 2000-922117		20000412
	EP 1666028 A1 20060607		EP 2005-27610		20000412
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, FI, CY				
	ES 2254164 T3 20060616		ES 2000-922117		20000412
	US 6518239 B1 20030211		US 2000-548759		20000413
	ZA 2002002855 A 20030821		ZA 2002-2855		20020411
	NO 2002001800 A 20020624		NO 2002-1800		20020417
	MX 2002PA04193 A 20021213		MX 2002-PA4193		20020426
	US 20030186894 A1 20031002		US 2002-313343		20021206
	US 6835372 B2 20041228				

	US 20050147567	A1	20050707	US 2004-985509	20041110
PRAI	US 1999-162451P	P	19991029		
	US 1999-164236P	P	19991108		
	US 1999-172769P	P	19991220		
	US 2000-178383P	P	20000127		
	US 2000-178415P	P	20000127		
	EP 2000-922117	A3	20000412		
	WO 2000-US9785	W	20000412		
	US 2000-548759	A1	20000413		
	US 2002-313343	A1	20021206		

AB The present invention provides a highly dispersible formulation comprising an active agent and a dipeptide or tripeptide comprising at least two leucyl residues. The composition of the invention possesses superior aerosol properties and is thus preferred for aerosolized administration to the lung. Also provided are a method for (i) increasing the dispersibility of an active-agent containing formulation for administration to the lung, and (ii) delivery of the composition to the lungs of a subject. The surface tension of several representative di- and tripeptides and proteins was determined and highly surface active peptides include dileucine and trileucine.

RE.CNT 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L2 ANSWER 4 OF 6 BIOSIS COPYRIGHT (c) 2008 The Thomson Corporation on STN
AN 2000:290609 BIOSIS
DN PREV200000290609
TI Method and apparatus for pulmonary administration of dry powder
alphan-antitrypsin.
AU Eljamal, Mohammed [Inventor, Reprint author]; Patton, John S. [Inventor]
CS Santa Clara, CA, USA
ASSIGNEE: Inhale Therapeutic Systems, Foster City, CA, USA
PI US 5993783 19991130
SO Official Gazette of the United States Patent and Trademark Office Patents,
(Nov. 30, 1999) Vol. 1228, No. 5. e-file.
CODEN: OGUPE7. ISSN: 0098-1133.
DT Patent
LA English
ED Entered STN: 6 Jul 2000
Last Updated on STN: 7 Jan 2002
AB Dry powders of alphan-antitrypsin are administered pulmonarily to patients
to treat, for example, certain types of emphysema. The dry powder
compositions may comprise aggregates of fine particles, which aggregates
are friable and break-up upon dispersion in a flowing gas stream.
Typically, the dispersed powders are captured in a chamber and
subsequently inhaled by a patient for pulmonary treatment of emphysema and
other conditions.

L2 ANSWER 5 OF 6 CAPLUS COPYRIGHT 2008 ACS on STN
AN 1998:478945 CAPLUS
DN 129:100052
TI Method and apparatus for pulmonary administration of dry powder
.alpha.1-antitrypsin
IN Eljamal, Mohammed; Patton, John S.
PA Inhale Therapeutic Systems, USA
SO U.S., 15 pp., Cont.-in-part of U.S. Ser. No. 423,515, abandoned.
CODEN: USXXAM
DT Patent
LA English
FAN.CNT 20

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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PI	US 5780014	A	19980714	US 1996-617512	19960313

EP 940154	A2	19990908	EP 1999-110369	19920702
EP 940154	B1	20070418		
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, MC, NL, SE				
EP 1693080	A2	20060823	EP 2006-9725	19920702
EP 1693080	A3	20070725		
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC				
AT 359842	T	20070515	AT 1999-110369	19920702
ES 2284226	T3	20071101	ES 1999-110369	19920702
US 6582728	B1	20030624	US 1995-423515	19950414
CA 2218208	A1	19961017	CA 1996-2218208	19960411
AU 9654825	A	19961030	AU 1996-54825	19960411
AU 703491	B2	19990325		
EP 866726	A1	19980930	EP 1996-911736	19960411
EP 866726	B1	20040303		
EP 866726	B2	20080109		
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, FI				
AT 260688	T	20040315	AT 1996-911736	19960411
ES 2217309	T3	20041101	ES 1996-911736	19960411
US 5993783	A	19991130	US 1998-114713	19980713
FRAI US 1995-423515	B2	19950414		
US 1991-724915	A	19910702		
EP 1992-914592	A3	19920702		
EP 1999-110369	A3	19920702		
US 1992-910048	A2	19920708		
US 1996-617512	A	19960313		
WO 1996-US5062	W	19960411		
AB	Methods are provided for administering .alpha.1-antitrypsin dry powder pulmonarily to a patient. In these methods, .alpha.1-antitrypsin is provided in a dry powder form which is aerosolized and administered to the patient. Apparatus are also provided for carrying out these methods. These methods and apparatus are may generally be used in the treatment of patients suffering from α 1-antitrypsin deficiency and the functional derangements of emphysema.			
RE.CNT 128	THERE ARE 128 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT			
L2	ANSWER 6 OF 6 CAPLUS COPYRIGHT 2008 ACS on STN			
AN	1986:223816 CAPLUS			
DN	104:223816			
OREF	104:35489a,35492a			
TI	Preparation of fat and protein from banked human milk: its use in feeding very-low-birth-weight infants			
AU	Hylmoe, P.; Polberger, S.; Axelsson, I.; Jakobsson, I.; Raeihae, N.			
CS	Nordreco AB, Bjuv, Swed.			
SO	Nestle Nutrition Workshop Series (1984), 5(Hum. Milk Banking), 55-61			
	CODEN: NNWSDT; ISSN: 0742-2806			
DT	Journal			
LA	English			
AB	Pooled human milk samples were heated to .apprx.50°, the cream was separated, and frozen and the skim milk (<0.5% fat) was subjected to ultrafiltration to remove lactose, water-soluble salts, and some low-mol.-weight proteins and to concentrate the protein fraction. The protein concentrate was freeze-dried and stored at -20°. When used to supplement mothers milk or bank milk (final protein concentration .apprx.2 g/100 mL and final fat concentration .apprx.5.5 g/100 mL) a slight increase in osmolality and Ca content was observed. The recovery of α -lactalbumin, lactoferrin, lysozyme [9001-63-2], and albumin in human milk protein supplement ranged from 60			

to 100% of that found in natural milk. The recovery of lactoferrin and IgA in the human milk protein concentrate was substantial and the powder addnl. contained .alpha.1-antitrypsin [9041-92-3], amylase [9000-92-4], and lipase [9001-62-1]. The use of the supplement in feeding very-low-birth-weight infants is discussed.